

CLAIMS:

1. A method for reducing or preventing apoptosis of differentiated cells selected from the list consisting of cardiac myocytes, pancreatic beta cells, endothelial cells and neuronal cells in the human or non-human mammal, which method comprises
5 administration, including acute administration, of an effective, non-toxic amount of a glucose uptake enhancer to a human or non-human mammal in need thereof.
2. A method for reducing or preventing apoptosis of cells induced by events selected from the list consisting of: ischaemic insult, serum deprivation, cytokine activation in the
10 human or non-human mammal, which method comprises administration, including acute administration, of an effective, non-toxic amount of a glucose uptake enhancer to a human or non-human mammal in need thereof.
3. A method for reducing post-ischaemic injury of the heart and/or improving the functional recovery of the heart following myocardial ischaemia which method comprises
15 administration of an effective, non-toxic amount of a glucose uptake enhancer to a human or non-human mammal in need thereof.
4. A method according to any one of claims 1 to 3, wherein the glucose uptake
20 enhancer is a thiazolidinedione.
5. A method according to claim 4, wherein the thiazolidinedione is Compound (I), or the tautomeric form thereof, or a pharmaceutically acceptable derivative thereof.
- 25 6. A method according to claim 4, wherein the thiazolidinedione is selected from:
(+) -5-[[4-[(3,4-dihydro-6-hydroxy-2, 5, 7, 8-tetramethyl-2H-1-benzopyran-2-yl)methoxy]phenyl]methyl]-2,4-thiazolidinedione (or troglitazone), 5-[4-[(1-methylcyclohexyl)methoxy]benzyl] thiazolidine-2,4-dione (or ciglitazone), 5-[4-[2-(5-ethylpyridin-2-yl)ethoxy]benzyl] thiazolidine-2,4-dione (or pioglitazone) or 5-[(2-benzyl-2,3-dihydrobenzopyran)-5-ylmethyl]thiazolidine-2,4-dione (or englitazone); or a
30 pharmaceutically acceptable derivative thereof.

7. A pharmaceutical composition comprising a glucose uptake enhancer, and a pharmaceutically acceptable carrier, wherein such composition is adapted for acute administration.

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